

Specifically, at the time that this application was filed, one skilled in the art of peptide synthesis would understand that the R-group side chain of the amino acid Glycine is a hydrogen atom. See, for example, Figs. 5-2 and 5-6 of Lehninger et al.; *Principles of Biochemistry, Second Edition*; Chapter 5 Amino Acids; Worth Publishers, Inc.; New York, NY; 1993; pp. 112 and 115 (copies attached). See also Panel 2-5 of Alberts et al.; *Molecular Biology of the Cell, Third Edition*; Garland Publishing, Inc.; New York, NY; 1994; pp. 56-57 (copy attached). See also the definition of amino acids of Coombs; *Dictionary of Biotechnology, Second Edition*; Stockton Press; New York, NY; 1992; p. 19 (copy attached). Thus, it would be clear to one skilled in the art that R2 of formulae I and II of claim 4 can be a hydrogen atom and an amino acid side chain.

For at least these reasons, claim 4 satisfies the requirements of 35 U.S.C. §112, second paragraph. Reconsideration and withdrawal of the rejection are respectfully requested.

II. Restriction Requirement

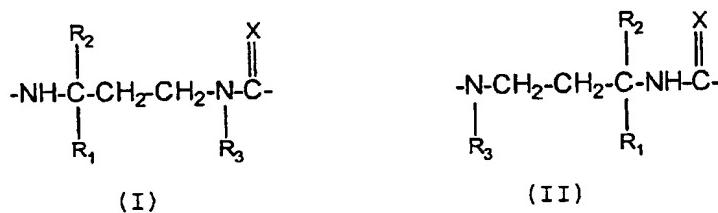
The Office Action makes the Restriction Requirement Final, and argues that the multiple individual compounds encompassed by formulae I and II cannot be a single special technical feature. Applicants respectfully assert that the Office Action misinterprets the PCT Unity of Invention Requirements, and applies the wrong standard in determining whether Unity of Invention exists between claims 1-18.

MPEP §1893.03(d) states that a group of inventions is considered linked to form a single inventive concept where there is a technical relationship among the inventions that involves at least one common or corresponding special technical feature. MPEP §1893.03(d) further states that the expression special technical feature is defined as meaning those technical features that define the contribution which each claimed invention, considered as a whole, makes over the prior art.

Under MPEP §1893.03(d), Unity of Invention exists between claims 1-18 where claims 1-18 are linked to form a single inventive concept. Claims 1-18 are linked to form a single inventive concept where claims 1-18 share a special technical feature, that is, where claims 1-18 share technical features that define the contribution that the claims as a whole make over the prior art.

All of claims 1-18 share the novel features of formulae I and II. Specifically, claim 1 recites the limitation:

at least one unit chosen from the B units of general formulae (I) and/or (II):



in which: R₁, R₂ and R₃ each independently of one another represent an amino acids side chain and may be identical or different, and X represents an oxygen or sulfur atom.

All of claims 2-18 ultimately depend from claim 1, and thus also include this feature.

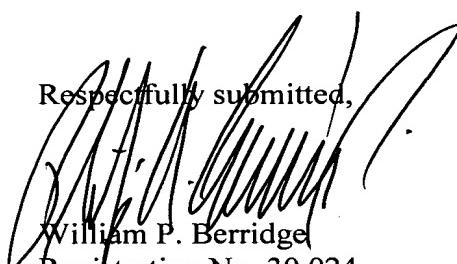
As indicated in the Office Action, claim 1 is allowed and this novel feature is not taught by the prior art. Because this novel feature defines a contribution that the claims as a whole make over the prior art, it is a special technical feature under MPEP §1893.03(d). Because claims 1-18 share this special technical feature, claims 1-18 are linked to form a single inventive concept under MPEP §1893.03(d). For these reasons, Unity of Invention exists between all of claims 1-18.

Because Unity of Invention exists between claims 1-18, the Restriction Requirement is improper and claims 12-17 should be rejoined to, and allowed with, claims 1-11 and 18. Reconsideration and withdrawal of the Restriction Requirement are respectfully requested.

III. Conclusion

In view of the foregoing, it is respectfully submitted that this application is in condition for allowance. Favorable reconsideration and prompt allowance of claims 1-18 are earnestly solicited.

Should the Examiner believe that anything further would be desirable in order to place this application in even better condition for allowance, the Examiner is invited to contact the undersigned at the telephone number set forth below.

Respectfully submitted,

William P. Berridge
Registration No. 30,024

Philip A. Caramanica, Jr.
Registration No. 51,528

WPB:PAC

Attachments:

Figs. 5-2 and 5-6 of Lehninger et al.; *Principles of Biochemistry, Second Edition*; Chapter 5 Amino Acids; Worth Publishers, Inc.; New York, NY; 1993; pp. 112 and 115.

Panel 2-5 of Alberts et al.; *Molecular Biology of the Cell, Third Edition*; Garland Publishing, Inc.; New York, NY; 1994; pp. 56-57.

Coombs; *Dictionary of Biotechnology, Second Edition*; Stockton Press; New York, NY; 1992; p. 19.

Date: June 30, 2005

OLIFF & BERRIDGE, PLC
P.O. Box 19928
Alexandria, Virginia 22320
Telephone: (703) 836-6400

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Amino Acids

Proteins can be reduced to their constituent amino acids by a variety of methods, and the earliest studies of proteins naturally focused on the free amino acids derived from them. The first amino acid to be discovered in proteins was asparagine, in 1806. The last of the 20 to be found, threonine, was not identified until 1938. All the amino acids have trivial or common names, in some cases derived from the source from which they were first isolated. Asparagine was first found in asparagus, as one might guess; glutamate was found in wheat gluten; tyrosine was first isolated from cheese (thus its name is derived from the Greek *tyros*, "cheese"); and glycine (Greek *glykos*, "sweet") was so named because of its sweet taste.

Amino Acids Have Common Structural Features

All of the 20 amino acids found in proteins have a carboxyl group and an amino group bonded to the same carbon atom (the α carbon) (Fig. 5-2). They differ from each other in their side chains, or R groups, which vary in structure, size, and electric charge, and influence the solubility of amino acids in water. When the R group contains additional carbons in a chain, they are designated β , γ , δ , ϵ , etc., proceeding out from the α carbon. The 20 amino acids of proteins are often referred to as the standard, primary, or normal amino acids, to distinguish them from amino acids within proteins that are modified after the proteins are synthesized, and from many other kinds of amino acids present in living organisms but not in proteins. The standard amino acids have been assigned three-letter abbreviations and one-letter symbols (Table 5-1), which are used as shorthand to indicate the composition and sequence of amino acids in proteins.

We note in Figure 5-2 that for all the standard amino acids except one (glycine) the α carbon is asymmetric, bonded to four different substituent groups: a carboxyl group, an amino group, an R group, and a hydrogen atom. The α -carbon atom is thus a **chiral center** (see Fig. 3-9). Because of the tetrahedral arrangement of the bonding orbitals around the α -carbon atom of amino acids, the four different substituent groups can occupy two different arrangements in space, which are nonsuperimposable mirror images of each other (Fig. 5-3). These two forms are called **enantiomers** or **stereoisomers** (see Fig. 3-9). All molecules with a chiral center are also **optically active**—i.e., they can rotate plane-polarized light, with the direction of the rotation differing for different stereoisomers.



Figure 5-2 General structure of the amino acids found in proteins. With the exception of the nature of the R group, this structure is common to all the α -amino acids. (Proline, because it is an imino acid, is an exceptional component of proteins.) The α carbon is shown in blue. R (in red) represents the R group or side chain, which is different in each amino acid. In all amino acids except glycine (shown for comparison) the α -carbon atom has four different substituent groups.

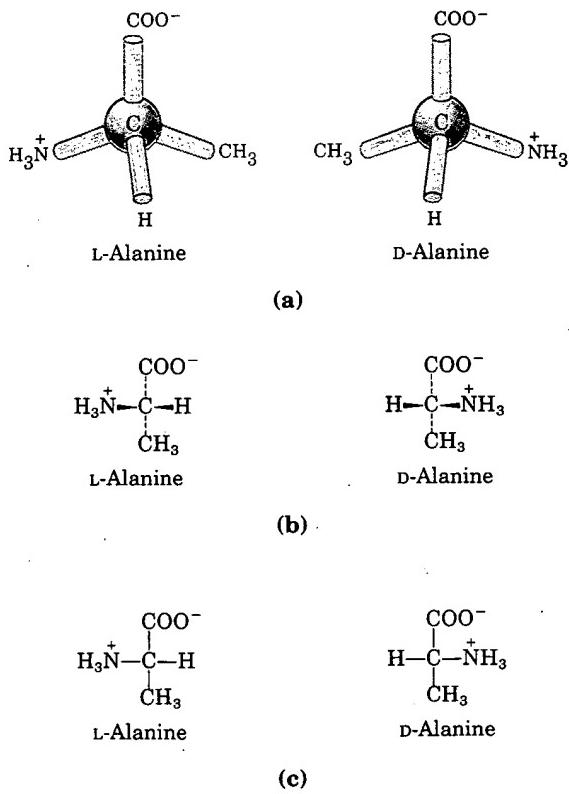


Figure 5-3 (a) The two stereoisomers of alanine. L- and D-alanine are nonsuperimposable mirror images of each other. (b, c) Two different conventions for showing the configurations in space of stereoisomers. In perspective formulas (b) the wedge-shaped bonds project out of the plane of the paper, the dashed bonds behind it. In projection formulas (c) the horizontal bonds are assumed to project out of the plane of the paper, the vertical bonds behind. However, projection formulas are often used casually without reference to stereochemical configuration.

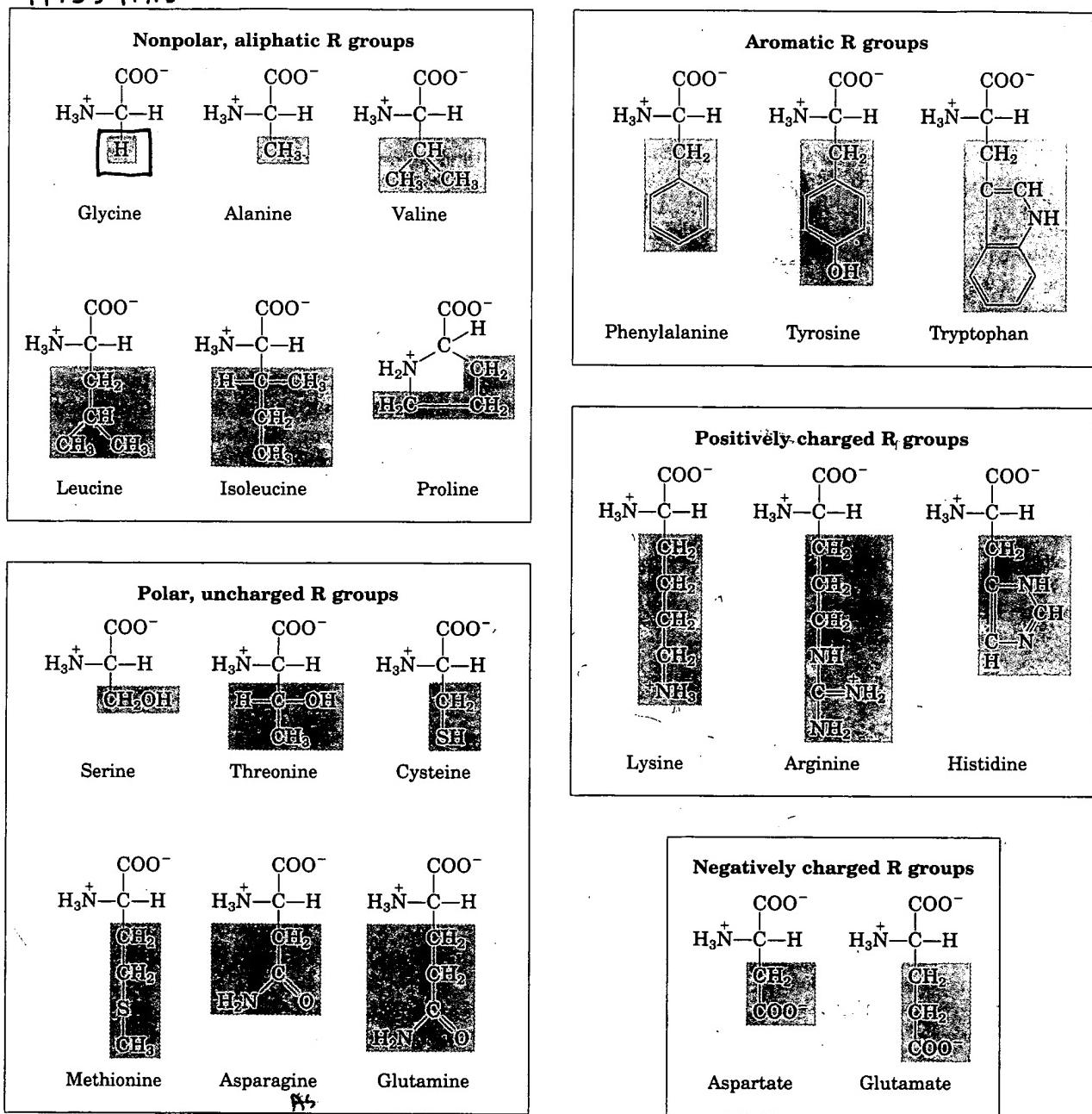
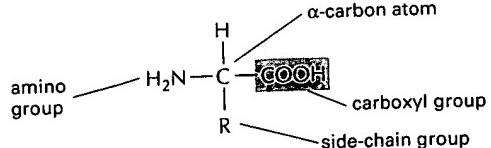


Figure 5–6 The 20 standard amino acids of proteins. They are shown with their amino and carboxyl groups ionized, as they would occur at pH 7.0. The portions in black are those common to all the amino acids; the portions shaded in red are the R groups.

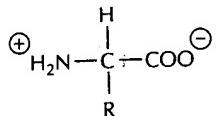
Nonpolar, Aliphatic R Groups The hydrocarbon R groups in this class of amino acids are nonpolar and hydrophobic (Fig. 5–6). The bulky side chains of alanine, valine, leucine, and isoleucine, with their distinctive shapes, are important in promoting hydrophobic interactions within protein structures. **Glycine** has the simplest amino acid structure. Where it is present in a protein, the minimal steric hindrance of the glycine side chain allows much more structural flexibility than the other amino acids. **Proline** represents the opposite structural extreme. The secondary amino (imino) group is held in a rigid conformation that reduces the structural flexibility of the protein at that point.

THE AMINO ACID

The general formula of an amino acid is

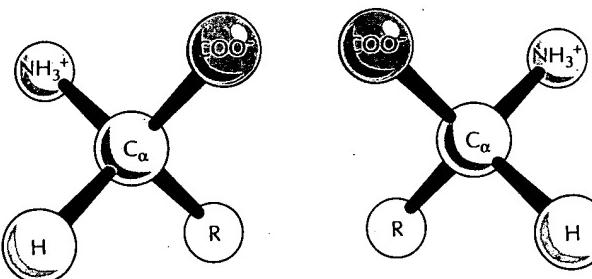


R is commonly one of 20 different side chains. At pH 7 both the amino and carboxyl groups are ionized.



OPTICAL ISOMERS

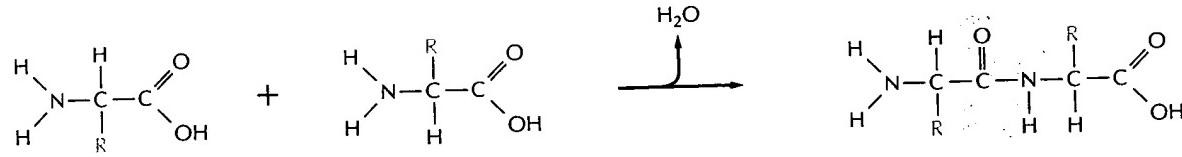
The α -carbon atom is asymmetric, which allows for two mirror image (or stereo-) isomers, D and L.



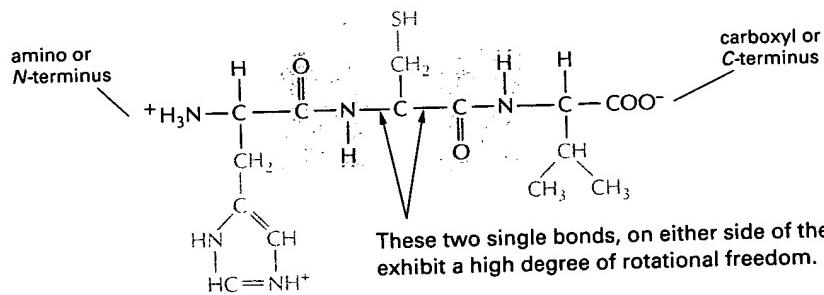
Proteins consist exclusively of L-amino acids.

PEPTIDE BONDS

Amino acids are commonly joined together by an amide linkage, called a peptide bond.



Proteins are long polymers of amino acids linked by peptide bonds, and they are always written with the N-terminus toward the left. The sequence of this tripeptide is His Cys Val.



These two single bonds, on either side of the rigid peptide unit, exhibit a high degree of rotational freedom.

FAMILIES OF AMINO ACIDS

The common amino acids are grouped according to whether their side chains are

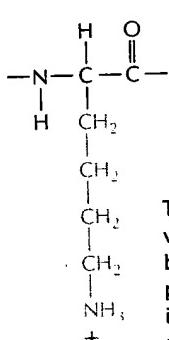
- acidic
- basic
- uncharged polar
- nonpolar

These 20 amino acids are given both three-letter and one-letter abbreviations.

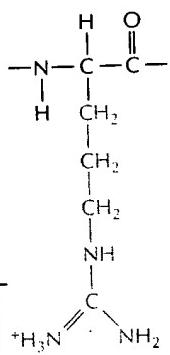
Thus: alanine = Ala = A

BASIC SIDE CHAINS

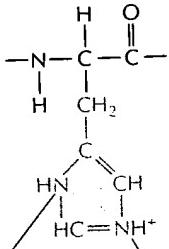
lysine
(Lys, or K)



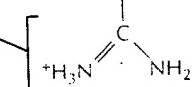
arginine
(Arg, or R)



histidine
(His, or H)



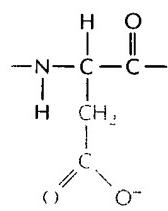
This group is very basic because its positive charge is stabilized by resonance.



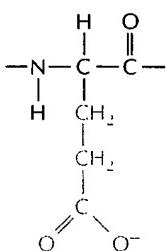
These nitrogens have a relatively weak affinity for an H^+ and are only partly positive at neutral pH.

ACIDIC SIDE CHAINS

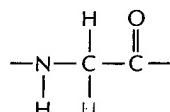
aspartic acid
(Asp, or D)



glutamic acid
(Glu, or E)

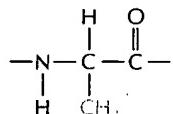


NONPOLAR SIDE CHAINS

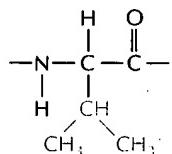


glycine
(Gly, or G)

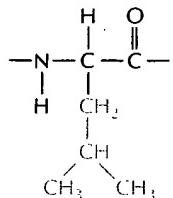
alanine
(Ala, or A)



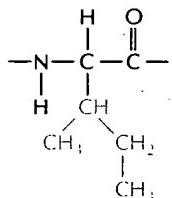
valine
(Val, or V)



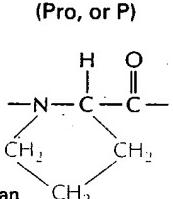
leucine
(Leu, or L)



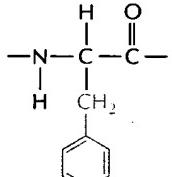
isoleucine
(Ileu, or I)



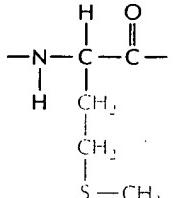
proline



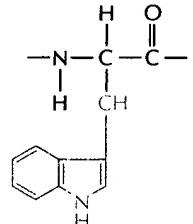
phenylalanine
(Phe, or F)



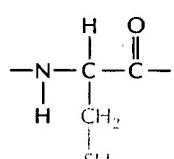
methionine
(Met, or M)



tryptophan
(Trp, or W)



cysteine
(Cys, or C)



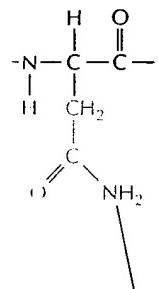
Amino acids with uncharged polar side chains are relatively hydrophilic and are usually on the outside of proteins, while the side chains on nonpolar amino acids tend to cluster together on the inside. Amino acids with basic or acidic side chains are very polar, and they are nearly always found on the outside of protein molecules.

The one letter code in alphabetical order:

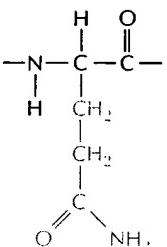
- Ala	G = Gly	M = Met	S = Ser
- Cys	H = His	N = Asn	T = Thr
- Asp	I = Ileu	P = Pro	V = Val
- Glu	K = Lys	Q = Gln	W = Trp
- Phe	L = Leu	R = Arg	Y = Tyr

UNCHARGED POLAR SIDE CHAINS

asparagine
(Asn, or N)



glutamine
(Gln, or Q)



Although the amide N is not charged at neutral pH, it is polar.

serine

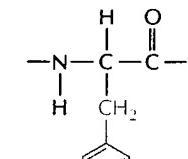
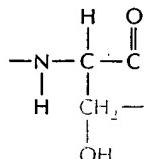
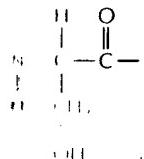
(Ser, or S)

threonine

(Thr, or T)

tyrosine

(Tyr, or Y)



OH group is polar.

Paired cysteines allow disulfide bonds to form in proteins.



5 amino acids per turn of the helix.
nation contributes to the secondary
ary structure of the protein.

interferon See leucocyte interferon.

ing double filter A wastewater
it process that consists of two bio-
ilters operating in series. Biomass
ates in the first filter as it consumes
the biochemical oxygen demand.
er of the filters is then reversed,
ie first filter becomes plugged with
which results in the rapid autolysis
umption of the starved biomass.

ion of generations A situation in
e life cycle of an organism contains
erent types of organism that differ
in appearance and mode of repro-
These differences, usually reflect
alternation between sexual and
generations, are common in plants
sistic animals. The sexual stage is
he gametophyte generation and the
stage the sporophyte. In most
e alternation of generations is as-
with alternations in diploid and
conditions and the intervention of
ind karyogamy. In some organisms
cycle may consist of more than two
ons which alternate regularly with
her.

A small sac such as occurs at the
of a bronchiole in the lungs or at
a duct in some glands.

abbreviation used to denote an
utant.

in An immunomodulator produced
'omyces.

nutation A class of suppressible
s that results in the creation of a
don in mRNA. This codon nor-
gnifies translation termination, so
ypeptide synthesis stops at the
ite. Such mutations can be sup-
n certain strains of *E. coli* possess-
NA with the AUC anticodon, and

hence inserting an amino acid at the UAG
site and permitting continued translation.

American Type Culture Collection
(ATCC) An organization holding a large
collection of microorganisms and cell lines,
including type specimens.

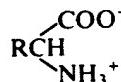
Ames test A test for potential mutagens and
frameshift carcinogens. Compounds are
screened for their ability to revert a series of
known frameshift mutants in the hisD gene
of *Salmonella typhimurium*. The reverted
cells can be recognized since they will grow
into colonies, which can be counted, on a
medium that lacks histidine.

amide An organic compound obtained by
replacing a hydroxyl group by an amino
group.

amido black A chemical stain used to
determine the position of proteins, includ-
ing products of antigen/antibody interac-
tion, on gels following chromatography or
electrophoresis.

amino acid analyser An automated ana-
lytical device designed to separate and
quantify individual amino acids from a
complex mixture which may have been
obtained from a protein hydrolysate or a
physiological fluid. In general, the analyser
consists of an automatic loading device to
which a large number of samples can be
added, a separation stage (usually based on
a chromatographic procedure) and a detec-
tion system (based on a colourimetric or flu-
orometric assay technique). Early systems
relied on two columns of ion exchange
resins which were sequentially eluted with
buffers of varying pH with the liquid stream
from the columns passing through a col-
ourimeter after having reacted with nin-
hydrin. The results were obtained as a trace
in which peaks of different height or area
indicated the presence of the various amino
acids. These could be quantified by using
standards to calibrate the machine. More
recent machines are fully automated single-
column liquid chromatographs, using
microprocessors for control and calibration
as well as calculation of the results which
may be printed directly.

amino acids Chemical compounds of the
following general formula



where R is a hydrogen atom (glycine) or any
of a number of different organic groups.
These compounds are zwitterions (dipolar
ions). Most amino acids found in biological
systems are present in the L-optical configura-
tion. Amino acids are the basic building
blocks of proteins, as well as participating
in central metabolism and contributing to
the synthesis of a variety of secondary pro-
ducts and biologically active molecules
including co-enzymes, hormones and neu-
rotransmitters. Several hundred different
amino acids are known, but only 20 are nor-
mally found in proteins. These may be clas-
sified on the basis of similarity in structure
or route of biosynthesis as shown in the
table. Other biologically important amino
acids include ornithine and citrulline which
are intermediates of the urea cycle, Y-ami-
nobutyric acid which functions as a neu-
rotransmitter, β -alanine which is a
precursor of pantothenic acid, and D-gluta-
mate which is found in bacterial cell walls.
Protein amino acids:

1. Hydrophobic

Alanine
Valine
Leucine
Isoleucine
Proline (or hydroxyproline)
Phenylalanine
Tryptophan
Methionine

2. Polar

Glycine
Serine
Threonine
Cysteine
Tyrosine
Asparagine
Glutamine

3. Acidic

Aspartic